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64 Tertiary aralkyl urethanes and isocyanates derived therefrom.

57 Production of tertiary aralkyl isocyanates, such as tetramethyl xylene diisocyanates, by thermal cracking of corresponding olefins and carbamic acid esters at moderate temperatures and in the presence of acid catalyst.

an alkali metal isocyanate (U.S. Patent 4,130,577) and by
reaction of the corresponding halide with isocyanic acid
(copending applications S.N. 331,696, filed December 17,
1981, and S.N. 335,945 filed March 8,
5 1982).

These various processes are disadvantageous for one
or more of a variety of reasons, such as that the materials
are difficult to handle or are corrosive, the yields are
poor, expensive reactants are required and the products
10 are difficult to recover.

It is an important object of this invention to produce
such tertiary aralkyl isocyanates utilizing non-corrosive,
low-cost starting materials in a simple process yielding
the desired isocyanates whereby they are readily recovered
15 and purified. In accordance with this invention the manu-
facture of a tertiary aralkyl isocyanate is by the pre-
paration of the corresponding tertiary aralkyl carbamic
acid ester of a lower aliphatic alcohol followed by the
thermal cracking of such urethane to form the isocyanate
20 and the free alcohol.

Since the esters of carbamic acid, such as methyl
carbamate, the esters of substituted carbamic acids, such
as the dimethyl ester of α, α, α' - tetramethylxylylene
dicarbamic acid, and the polymers of poly isocyanates with
25 polyols are all properly called urethanes, for the sake of
clarity, herein an unsubstituted carbamic acid ester will
be called "carbamate" or "carbamic acid ester"; a sub-
stituted carbamic acid ester will be called "urethane",
"monourethane", "diurethane", "polyurethane" or "urethane
30 ester"; and the polymers will be called "urethane polymers".

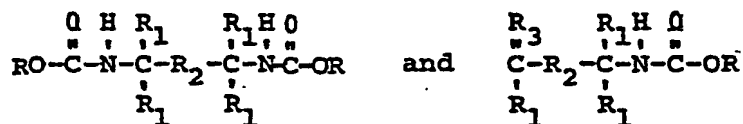
Urethanes which are useful in forming tertiary aralkyl
isocyanates by thermal cracking in accordance with this
invention are generally designated by the formulae:

TERTIARY ARALKYL URETHANES AND
ISOCYANATES DERIVED THEREFROM

This invention relates to the manufacture of isocyanates and in particular provides a new route for the synthesis of tertiary aralkyl isocyanates and also provides a valuable class of urethanes useful in the production of tertiary aralkyl isocyanates.

Isocyanates having an isocyanato group attached to a tertiary aliphatic carbon atom also having aromatic substituents, for example, meta- and para- tetramethylylene diisocyanates, are particularly useful for reaction with a wide variety of polyols to give urethane polymers which can be either rigid or flexible and which can be endowed with a wide variety of properties. Such urethane polymers can be formed into rigid and flexible foamed articles, sheets, high density sheets and articles of various shapes. The light stability of these urethane polymers makes them extremely useful in surface coatings and other applications where light stability is desirable, e.g. light stable RIM elastomers.

Such tertiary aralkyl isocyanates have been prepared by a variety of different reactions, such as by the phosphorylation of corresponding organic amines, by reaction of corresponding olefins with isocyanic acid (U.S. Patent 3,290,350), by reaction of the corresponding halides with



wherein R is an alkyl radical having from 1 to 18 or more carbon atoms; R_1 is an alkyl radical having from 1 to 3 carbon atoms; and R_2 represents an aromatic hydrocarbon moiety such as phenyl, biphenyl and naphthalyl and such an aromatic hydrocarbon moiety having substituents including halogen atoms, methyl and methoxy groups and substituents such as:



and wherein R_3 is an alkylidene group having from one to three carbon atoms.

These urethanes, which themselves are an important part of this invention, are thus tertiary aralkyl urethanes in which R is a radical derived from an aliphatic alcohol, such as methanol, ethanol, isopropanol, 2-ethyl-hexanol-1 and n-octadecanol. Such alcohol is split off on thermal cracking of the urethane ester and can usually be recovered and recycled as discussed below.

The tertiary aralkyl urethane esters of this invention can be prepared, for example, by reaction of the carbamic acid ester of a lower aliphatic alcohol and an olefin, as for example, is disclosed in Belgian Patent No. 625,748, granted December 12, 1962. As set forth in that patent the reaction of the olefin and carbamate generally proceeds at moderate temperatures in the presence of an acid cat-

alyst.

The urethane esters of this invention are converted to the corresponding tertiary aralkyl isocyanates by thermal cracking. Yields are good, and alcohol recovery and removal are particularly high where the aliphatic alcohol group of the initial carbamic acid ester is methanol or other lower, preferably normal alcohol.

In accordance with this invention the preparation of the tertiary aralkyl urethane ester takes place at temperatures from 40°C up to 150°C in the presence of an acid catalyst, such as sulfuric acid, toluene sulfonic acid, dodecyl benzene sulfonic acid, hydrocarbon sulfate esters, hydrochloric acid, boron trifluoride and other Lewis and Bronstead acids. The reaction can take place in the absence of solvent or in the presence of solvents, such as methylene dichloride, toluene, xylene, chlorobenzene, and so forth.

Generally, the proportion of carbamic acid ester to olefin is in excess of stoichiometric. Preferably the carbamate is in substantial excess and functions as solvent and catalyst moderator as well as reactant. It is preferred in accordance with this invention to use from 50% to 800% stoichiometric excess of carbamate.

The amount of catalyst required to promote the addition of tertiary aralkyl olefin and carbamic acid ester is not critical and can be varied widely. Where substantial excess of carbamic acid ester is utilized the amount of catalyst, based on the olefin, is typically 0.01 to 10 mole% and preferably about 2 mole %.

Preferably the carbamate is heated mildly to maintain it molten, from 40° to 150°C being preferred. The catalyst is dissolved in the molten carbamate, and the olefin is then slowly added. When the reaction is complete, the mixture is treated to remove or neutralize the catalyst. Unreacted

carbamate ester is then removed by distillation.

Tertiary aralkyl urethane esters in accordance with this invention form the corresponding isocyanate by thermal cracking while splitting off the alkanol. In many cases the alcohol can usefully be recycled by reaction with urea or isocyanic acid (HNCO) to form the starting carbamate ester. In addition to their utility in the formation of the corresponding isocyanates the tertiary aralkyl urethane esters of this invention may also be useful for other purposes. For example, some of these compounds are quite similar to known herbicides, and herbicidal activity is anticipated. The urethane esters of higher alkanols can also be used as blocked isocyanates in powder coatings and the like.

In cracking the urethane esters according to this invention to form the corresponding isocyanates the catalyst must be removed or neutralized for example, with calcium oxide, sodium carbonate, sodium hydroxide and the like, which is followed by cracking of the urethane ester either neat or in high boiling solvents, such as hexadecane, diphenyl ether, diisopropyl naphthalene and the like. Cracking takes place at temperatures on the order of 150° to 350°C. splitting off the alkanol to yield the corresponding isocyanate.

This invention has particular application in the production of tertiary aralkyl polyisocyanates from polyolefins and carbamic acid esters by addition of the reactants to form the corresponding polyurethane esters followed by thermal cracking of the polyurethane esters to the corresponding polyisocyanates. Polyolefin reactants which are contemplated include diisopropenyl benzenes (DIPEB), triisopropenylbenzene and diisopropenyl naphthalenes, all of which yield highly desirable polyisocyanates. The addition reaction can also be utilized with diolefins, such as a diisopropenylbenzene, to favor the production of a monourethane, such as isopropenyl- α,α -dimethylbenzyl urethane (TMU), which can be thermally cracked to the corresponding olefinic

monoisocyanate, such as isopropenyl- α,α -dimethylbenzyl isocyanate (TMI).

The thermal cracking of tertiary aralkyl polyurethanes derived by an acid promoted addition reaction to form the corresponding isocyanates is particularly surprising in view of the instability of such urethanes in the presence of even a small amount of acid. Polyurethanes, such as tetramethylxylylene diurethane (TMXDU), crack extensively in the presence of traces of acid to form monoisocyanates and other undesired products. It is, therefore, also a feature of the invention that the urethane esters are thermally cracked only after complete neutralization of the acid catalyst. By "complete neutralization" it should be understood that the amount of alkali or alkaline material, such as calcium oxide, sodium hydroxide, sodium carbonate or the like, is not just in excess of the stoichiometric quantity but is sufficient to bring the pH of the mixture substantially to 7 or higher. Neutralization should be carried out before recovery steps, such as removal of the excess carbamic acid ester after which the urethane ester can be cracked.

The following Examples illustrate the preparation of urethane esters in accordance with this invention and the utilization of such esters in the preparation of isocyanates useful in preparation of polyurethanes.

EXAMPLE 1.

A mixture containing 28.76g (383.10 mmoles) of methyl carbamate, 8.8ml of 1.08M p-toluenesulfonic acid (9.50 mmoles) in toluene solution and 40ml of methylene chloride was stirred at 45°C until all solid was dissolved. Meta-diisopropenyl benzene (m-DIPEB) (15.19g, 96.1 mmoles) was then added dropwise into the

solution. After stirring at 45°C for 22 hours, the reaction mixture was cooled to room temperature, washed with 10% aqueous sodium carbonate solution, and the organic layer dried over sodium sulfate. Analysis by GLC (heneicosane, $C_{21}H_{44}$, internal standard) indicates a 67% yield of $\alpha, \alpha, \alpha', \alpha'$ -tetramethyl-meta-xylylene diurethane (m-TMXDU) and a 20% yield of the methyl ester of m-isopropenyl - α, α -dimethyl-benzyl urethane (TMU).

- 10 Concentration of the resulting methylene chloride reaction mixture was carried out at reduced pressure. An equal volume of hexane was then added to precipitate the dimethyl ester m-TMXDU, giving a white solid which was collected by filtration (mp 128-130°C; yield: 44%
- 15 m-TMU). The remainder of the m-TMXDU in the methylene chloride-hexane solution could be isolated by further evaporation of methylene chloride solvent, followed by the addition of more hexane to the oily residue.

EXAMPLE 2

- 20 To a flask containing 59.30g (789.92 mmols) of molten methyl carbamate (85°C oil bath) was added 260 mg (2.6 mmole) of conc. sulfuric acid. The temperature of the oil bath was then lowered to 60°C, and 14.79g (93.61 mmols) of m-DIPEB added dropwise to the reaction mixture. The reaction mixture was stirred 60°C for 4.5 hours, and
- 25 500 mg of anhydrous sodium carbonate added to neutralize the acid catalyst. The mixture was stirred at 60°C for another 30 minutes and then distilled under the vacuum (95°/30mm Hg) to recover unreacted methyl carbamate (44.05g; > 90% recovery). GLC analyses of the white
- 30 residue showed that m-TMXDU and m-TMU were formed in 65% and 24% yields, respectively. The solid was then mixed with 50g of hexadecane and 1.66g of heneicosane

(GLC internal standard), and the mixture cracked at 265°C for 4 hours. GLC analyses indicated that $\alpha, \alpha, \alpha', \alpha'$ -tetramethyl meta-xylene diisocyanate (m-TMXDI) (45%), meta-isopropenyldimethylbenzene isocyanate (m-TMI) (10%) and $\alpha, \alpha, \alpha', \alpha'$ -tetramethyl-meta-xylene mono-urethane monoisocyanate (m-TMXUI) (3%) were formed based on the m-DIPEB.

EXAMPLE 3

To a flask containing 59.30g (787.92 mmoles) of molten methyl carbamate (85°C oil bath) was added 260 mg (2.6 mmoles) of concentrated sulfuric acid. The temperature of the reaction mixture was then lowered to 60°C and 14.79g (93.61 mmoles) of p-DIPEB added dropwise to the reaction mixture. The reaction mixture was stirred for 4.5 hours at 60°C, and 1g CaO added to neutralize the acid catalyst. The reaction mixture was stirred for another 30 minutes at 60°C and distilled under vacuum (95°/30mm Hg) to recover unreacted methyl carbamate (45.0g; > 90% recovery). GLC analysis of the residue showed that 69% p-TMXDU and 21% p-TMU had formed.

The resultant mixture was then cracked by the treatment described for the corresponding meta-compounds in Example 2. GLC analysis gave 62% p-TMXDI and 16% p-TMI based on the p-DIPEB.

EXAMPLE 4

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By way of further examples of the thermal cracking of diurethane, a mixture containing 25.15g of p-TMXDU (81.6 mmoles), 100g of hexadecane and 14.25 g of heneicosane (an inert internal standard) was heated with stirring in a 265°C oil bath in a slow stream of nitrogen. Methanol evolved was collected in a dry-ice acetone cold trap. Heating

continued for 4 hours, and the reaction mixture analyzed with GLC and IR after cooling to room temperature. Both GLC and IR analysis confirmed the formation of p-TMXDI, p-TMXUI (mono-urethane-monoisocyanate) and p-TMI (mono-isopropenyl-monoisocyanate). GLC (internal standard method) indicated yields were 68%, 20% and 6%, respectively. The liquid collected in the cold trap weighed 4.57g, and was mostly methanol, as indicated by IR analyses.

EXAMPLES 5-27

10 Although the preceding examples have been confined to urethanes produced by reaction of tertiary aralkyl olefins with methyl carbamate, urethanes which are esters of other lower alkanols, particularly straight chain alkanols, are useful in preparing isocyanates by thermal
15 cracking following the procedures generally outlined above. A number of such urethane esters have been prepared and are described in TABLE I.

T A B L E I

EX. NO.	URETHANE ESTER	ALKANOL RESIDUE	APPEARANCE
5	$\text{ROOCHN}-\text{C}_6\text{H}_4-\text{NHCOOR}$	methyl	white solid
6	"	ethyl	white fluffynedlea
7	"	n-propyl	white needles
8	"	iso-propyl	white powder
9	"	n-butyl	white powder
10	"	2-ethylhexyl	white powder
11	"	cyclohexyl	white powder
12	$\text{ROOCHN}-\text{C}_6\text{H}_4-\text{NHCOOR}$	methyl	white crystalline powder
13	"	ethyl	white needles (fine)
14	"	n-propyl	fine white needles
15	"	iso-propyl	white powder
16	"	n-butyl	white crystals solid

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TABLE I (Continued)

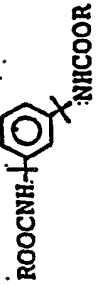

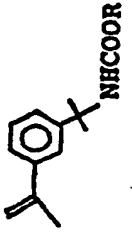
EX. NO.	URETHANE ESTER	ALKANOL RESIDUE	APPEARANCE
17		benzyl	white solid
18	" " "	n-octadecyl	white solid
19		methyl	white solid
20		methyl	oil
21	" " "	ethyl	oil
22	" " "	n-propyl	oil
23	" " "	iso-propyl	oil
24	" " "	n-butyl	oil
25	" " "	benzyl	oil
26	" " "	n-octadecyl	white solid

TABLE I (Continued)APPEARANCE

white solid

ALKANOL RESIDUE

methyl

URETHANE ESTEREX. NO.

27

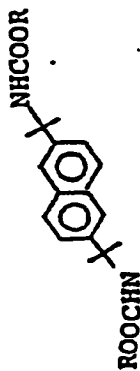


TABLE I (Continued)

EX. NO.	M.P., °C	NMR δ-VALUE (CDCl ₃) (60-MHz)
5	182-184	<u>1.6</u> (12H, s, CH ₃); <u>3.5</u> (6H, s, O-CH ₃); <u>5.1</u> (2H, bs, NH); <u>7.25</u> (4H, s, aro)
6	159-160	<u>1.15</u> (6H, t, J=7Hz); <u>1.64</u> (12H, s); <u>3.98</u> (4H, q, J=7Hz); <u>5.05</u> (2H, bs); <u>7.30</u> (4H, s)
7	170-173	<u>0.83</u> (6H, t, J=7Hz); <u>1.2-1.8</u> (4H, m, J=7Hz); <u>1.65</u> (12H, s); <u>3.88</u> (4H, t, J=7Hz); <u>5.03</u> (2H, bs); <u>7.25</u> (4H, s)
8	196-197	<u>1.12</u> (12H, d, J=6.5Hz); <u>1.65</u> (12H, s); <u>4.77</u> (2H, septet, J=6.5Hz); <u>4.97</u> (2H, bs); <u>7.30</u> (4H, s)
9	132-133.5	<u>0.86</u> (6H, t, J=6Hz); <u>1.0-1.8</u> (8H, m); <u>1.65</u> (12H, s); <u>3.91</u> (4H, t, J=6Hz); <u>5.00</u> (2H, bs); <u>7.22</u> (4H, s)
10	97-98	<u>0.86-1.8</u> (30H, m); <u>1.63</u> (12H, s); <u>3.86</u> (4H, d, J=4.5Hz); <u>5.00</u> (2H, bs); <u>7.25</u> (4H, s)
11	202-204	<u>1.0-1.9</u> (22H, m); <u>1.63</u> (12H, s); <u>4.5</u> (2H, m); <u>4.93</u> (2H, bs); <u>7.23</u> (4H, s)
12	130-132	<u>1.66</u> (12H, s); <u>3.55</u> (6H, s); <u>5.10</u> (2H, bs); <u>7.20</u> (3H, m); <u>7.33</u> (1H, m)
13	101-103	<u>1.14</u> (6H, t, J=7Hz); <u>1.64</u> (12H, s); <u>3.97</u> (4H, q, J=7Hz); <u>5.02</u> (2H, bs); <u>7.15</u> (3H, s); <u>7.30</u> (1H, m)
14	94-95.5	<u>0.83</u> (6H, t, J=6.5Hz); <u>1.2-1.8</u> (4H, m); <u>1.63</u> (12H, s); <u>3.85</u> (4H, t, J=6.5Hz); <u>4.98</u> (2H, bs); <u>7.13</u> (3H, m); <u>7.28</u> (1H, m)

TABLE I (Continued)

EX. NO.	M.P., °C	NMR δ-VALUE (CDCl ₃) (60-MHz)
15	105-107.5	1.12 (12H, d, J=6Hz), 1.63 (12H, s), 4.73 (2H, septet), 4.93 (2H, bs), 7.12 (3H, m), 7.27 (1H, m)
16	64-66.5	0.86 (6H, t, J=6Hz), 1.1-1.8 (8H, m), 1.64 (12H, s), 3.90 (4H, t, J=6Hz), 4.98 (2H, bs), 7.17 (3H, m), 7.27 (1H, m)
17	82-84	
18	47-49	
19	56-57	1.65 (6H, s), 2.12 (3H, dd, J ₁ =1.6, J ₂ =0.8Hz), 3.53 (3H, s), 4.97 (1H, dq, J ₁ =J ₂ =1.6Hz), 5.05 (1H, bs), 5.25 (1H, dq, J ₁ = 1.6, J ₂ =0.8), 7.25 (4H, s)
20	bp 115 at 1mm	1.55 (6H, s), 2.05 (3H, dd), 3.45 (3H, s), 4.95 (1H, dq), 5.25 (1H, bs), 5.65 (1H, dq), 7.16 (3H, m), 7.40 (1H, m)
21	---	1.13 (3H, t, J=7Hz), 1.64 (6H, s), 2.10 (3H, dd), 3.95 (2H, q, J=7Hz), 4.97 (1H, dq), 5.03 (1H, bs), 5.20 (1H, dq), 7.13 (3H, m), 7.32 (1H, m)
22	---	0.82 (3H, t, J=7Hz), 1.2-1.8 (2H, m), 1.65 (6H, s), 2.10 3H, dd), 3.85 (2H, t, J=7Hz), 4.97 (1H, dq), 5.02 (1H, bs), 5.20 (1H, dq), 7.14 (3H, m), 7.30 (1H, m)

TABLE 1 (Continued)

EX. NO.	M.P., °C	NMR δ-VALUE (CDCl ₃) (60-MHz)
23	---	1.12 (6H, d, J=6Hz), 1.63 (6H, s), 2.12 (3H, dd, J=1.6, J ₂ =0.8), 4.74 (1H, septet), 4.98 (1H, dq, J ₁ =J ₂ =1.6), 5.03 (1H, bs), 5.23 (1H, dq, J ₁ =1.6, J ₂ =0.8), 7.17 (3H, m), 7.35 (1H, m)
24	---	0.85 (3H, t), 1.1-1.8 (4H, m), 1.65 (6H, s), 2.12 (3H, dd), 3.90 (2H, t), 4.97 (1H, dq), 5.10 (1H, bs), 5.23 (1H, dq), 7.17 (3H, m), 7.23 (1H, m)
25	---	
26	78-81	
27	177-178	7.50 (bm, 3, Ar), 5.15 (bs, 1, NH), 3.52 (s, 3, OCH ₃), 1.70 (s, 6, CH ₃)

In Table I the urethane esters of alkanols other than methanol were prepared by reaction of the alkanol and the corresponding isocyanate, rather than by the addition reaction of carbamate ester and olefin. The addition reaction can, however, be employed to prepare these urethanes also. Yields of diurethanes are not as good as yields of monourethanes. The yields also are reduced as higher alkanols are used. Yields are generally better with normal alkanol esters of carbamic acid.

In a commercial processing arrangement the olefinic monourethane product TMU can be recycled with DIPEB in addition reaction with methyl carbamate. The two urethanes TMXDU and TMU are difficult to separate, however; and recycle of the TMU is not practicable. The thermally cracked product of the monourethanes, the isopropenyl monoisocyanate TMI, moreover, can not be recycled directly as it will not react with methyl carbamate under the reaction conditions for DIPEB and carbamate ester. Nevertheless, where this product is not desired for other purposes it can be reacted with methanol to reform the isopropenyl-monourethane TMU for recycle with DIPEB. The monourethane-monoisocyanate product of the thermal cracking stage can, of course, be recycled for additional thermal cracking. The following are Examples:

EXAMPLE 28

m-DIPEB was slowly added to molten mixture of methyl carbamate (mc) and concentrated sulfuric acid maintained at 60°C (mole ratio of m-DIPEB: MC: H_2SO_4 = 1:8:0.025, respectively). After the addition of DIPEB, the reaction mixture was stirred for 4-5 hours, and 300% mole excess solid sodium carbonate relative to sulfuric acid was added to basify the mixture. The mixture was stirred at 60°C for

a half hour more and the reaction mixture vacuum distilled (approx. 80°/20 mm.) to recover unreacted methyl carbamate (>90% recovered). To the distillate residue which contained m-TMU (24% yield) and m-TMXDU (65% yield) was
5 added 10% by weight of finely divided CaO. The mixture was heated at 200° at 20 mmHg. for 90 minutes to crack the urethanes to m-TMXDI and m-TMI. At the end of the cracking cycle, the vacuum was decreased to 10mm and m-TMXDI and m-TMI collected by fractional distillation.
10 The m-TMI collected was recycled as described above, by first converting it to m-TMU by reaction with methanol.

EXAMPLE 29

A solution containing 34.2g (170 mmoles) of m-TMI, 100ml of methanol and 50mg of dibutyltin dilaurate was
15 heated with stirring in a 70°C oil bath for 3 hours. The reaction mixture was then cooled to room temperature, and the excess methanol was removed at reduced pressure. The residue, an oil, was distilled under vacuum to yield 34.2 grams of m-TMU (bp 115°C, 1mm Hg).

20

EXAMPLE 30

m-TMU 5g (22 mmoles) was slowly added to a molten mixture of methyl carbamate (13.2g; 176 mmoles) containing 0.26g (2.6 mmoles) of concentrated sulfuric acid at 70°C. The mixture was heated for three hours at 70°, and analyzed
25 by GLC in the usual manner. Analysis showed an 84% yield of m-TMXDU with 15% m-TMU unreacted.

EXAMPLE 31

To a molten mixture of 20.0g (266.4 mmoles) of methyl

carbamate and 100mg (1 mmole) of conc. sulfuric acid maintained at 65°C (oil bath) was added 5.2g (32.9 mmole) of m-DIPEB and 2.4g (10.3 mmole) of m-TMU. After the addition was completed, the reaction was stirred at 65°C for 3 hours. and analyzed by GLC. Analysis indicated that the area percent ratio of m-TMXDU to m-TMU was the same as that observed in the m-DIPEB-methyl carbamate reaction (4~5:1). The yields of m-TMXDU and m-TMU were 72 and 18% respectively.

EXAMPLE 32

A processing arrangement for production of TMXDI based on the preceding examples, particularly 29-31, is illustrated in the accompanying drawing which is a flow diagram.

As shown in the drawing, there are four basic steps involved in the production of TMXDI from DIPEB and the methyl ester of carbamic acid. These are the addition reaction of the carbamate and diolefin, indicated by the reference numeral 10; neutralization of the acid catalyst with base, indicated by the reference numeral 20; stripping of the methyl carbamate, indicated by the reference numeral 30; and thermal cracking of the resultant urethanes to TMXDI and other products, indicated by the reference numeral 40.

In this process the two basic ingredients, DIPEB and methyl carbamate are introduced at addition step 10, as indicated by flow lines 11 and 12. Catalyst, such as sulfuric acid, is also introduced as indicated by the reference numeral 13. Recycled TMU subsequently formed during the process is, as shown by flow line 15, is also introduced at addition step 10.

The product of olefin addition step 10 is admixed with base, such as sodium carbonate in the neutralization

step 20, as indicated by flow lines 21 and 22 respectively. The neutralized material, as indicated by flow line 31, is then stripped of methyl carbamate at step 30 to recover methyl carbamate which via line 14 is recycled to addition step 10 with make-up methyl carbamate indicated by line 16. After stripping the methyl carbamate, the product as indicated by flow line 41 consisting basically of TMXDU, TMU and polymerized material, is then treated at cracking step 40. TMI and methyl alcohol are recovered in cracking step 40 as indicated by the flow lines 51 and 52, respectively. The desired product TMXDI is recovered from cracking step 40 as indicated by flow line 43, and still bottoms from cracking step 40 are indicated by flow line 44.

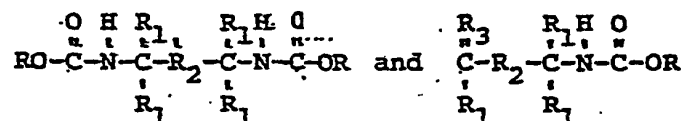
To the extent by-product TMI can not be used as such it is reacted with the recovered methanol by a process 50 such as described in Example 29 to convert the olefinic TMI to TMU, which then is recycled to the addition step, as indicated by flow line 15, for further reaction with methyl carbamate.

An important aspect of the recycle of TMU is that the product distribution in olefin addition step 10 is not affected by the presence of the added olefinic TMU. Thus the recycle of TMI as TMU greatly improves the efficiency in conversion of DIPEB to TMXDI.

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THE CLAIMS

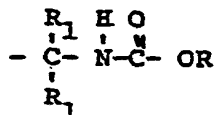
1. Urethanes selected from urethanes having the formulae:



wherein R is an alkyl radical having from 1 to 18 carbon atoms;

R₁ is an alkyl radical having from 1 to 3 carbon atoms;

R₂ represents an aromatic hydrocarbon moiety selected from phenyl, biphenyl and naphthyl groups and such groups having halo, methyl and methoxy substituents and substituents of the formula:



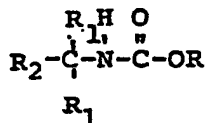
and wherein R₃ is an alkylidene radical having from 1 to 3 carbon atoms.

2. The alkyl esters of $\alpha, \alpha, \alpha', \alpha'$ -tetramethyl-meta-xylylene dicarbamic acid.

3. The dialkyl esters of $\alpha, \alpha, \alpha', \alpha'$ -tetramethyl-para-xylylene dicarbamic acid.

4. The alkyl esters of isopropenyl cumyl carbamic acid.

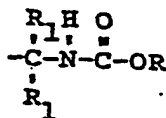
5. In a process for production of urethanes of the general formula:



where R is an alkyl radical derived from a lower alkanol;

R₁ is an alkyl radical having from 1 to 3 carbon atoms; and

R₂ represents an aromatic hydrocarbon moiety selected from phenyl, biphenyl and naphthalyl groups and such groups having halo, methyl and methoxy substituents and substituents of the formula:



wherein R and R₁ have the same significance, by addition of the corresponding olefin and a carbamic acid ester of a lower alkanol in the presence of moderate heat and an acid catalyst, the improvement which comprises forming a mixture consisting essentially of said catalyst and said carbamic acid ester at a temperature between 40° and 150°C at which said carbamic acid ester is molten and thereafter adding said olefin to said mixture to react said carbamic acid ester and olefin to form said urethane.

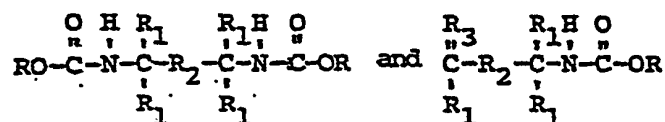
6. The improvement according to claim 5 in which said olefin is added to said mixture dropwise, the olefin is a diisopropenyl benzene and the carbamic acid ester is methyl carbamate.

7. The improvement according to claim 5, or 6 in which said mixture and added olefin are neutralized after formation of said urethane.

8. The improvement according to claim 5 in which said carbamic acid ester is in excess of the amount of ole-

fin required to form said urethane, in which said mixture and added olefin are neutralized after formation of said urethane, and in which excess, unreacted carbamic acid ester is removed after neutralization.

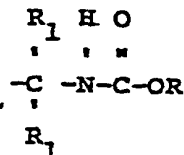
9. A process for the production of tertiary ar-alkyl isocyanates which comprises heating a urethane of the formulae



wherein R is an alkyl radical derived from a lower alkanol;

R₁ is an alkyl radical having from 1 to 3 carbon atoms; and

R₂ is an aromatic hydrocarbon moiety selected from phenyl, biphenyl and naphthalyl groups and such groups having halo, methyl and methoxy substituents and substituents of the formula:



and wherein R₃ is an alkylidene group having from one to three carbon atoms, said urethane being heated at 200°-300° for a sufficient time to split off the alkanol of the formula ROH and form the isocyanate corresponding to said urethane.

10. A process for production of α,α,α',α'-tetramethyl xylylene diisocyanates which comprises:

(a) reacting a diisopropenyl benzene and isopropenyl-, -dimethylbenzyl urethane with an excess of methyl carbamate in the presence of moderate heat and an acid catalyst to convert the olefins to urethanes,

(b) neutralizing said catalyst in the resultant mixture,

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(c) stripping unreacted methyl carbamate from the neutralized mixture,

(d) thermally cracking the stripped mixture to convert urethanes to isocyanates whereby a mixture including tetramethyl xylylene diisocyanate and isopropenyl dimethylbenzyl isocyanate is obtained,

(e) recovering methanol from said mixture during thermal cracking,

(f) separating isopropenyl-dimethylbenzyl isocyanate and tetramethyl xylylene diisocyanate from the cracked mixture,

(g) reacting said isopropenyl dimethylbenzyl isocyanate and methanol to form isopropenyl-dimethylbenzyl urethane and,

(h) recycling said isopropenyl-dimethylbenzyl urethane so formed and said stripped methyl carbamate to said reaction with diisopropenyl benzene.





European Patent
Office

EUROPEAN SEARCH REPORT

0101832

Application number

EP 83 10 6424

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 7)
A	DE-C-1 157 598 (BAYER AG) * Example 2 * & BE-A-625 748 (Cat. D)		C 07 C 125/073 C 07 C 125/065 C 07 C 119/042 C 07 C 118/00
D,A	US-A-4 130 577 (SHOWA DENKO) * Abstract *		
D,A	US-A-3 290 350 (E.I. DU PONT DE NEMOURS) * Example 6 *		
			TECHNICAL FIELDS SEARCHED (Int. Cl. 7)
			C 07 C 118/00 C 07 C 119/042 C 07 C 125/00
The present search report has been drawn up for all claims			
Place of search BERLIN		Date of completion of the search 28-09-1983	Examiner BREW C.H.
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	

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